

ANTIMICROBIAL CONTAINING SOLVENTLESS

HOT MELT ADHESIVE COMPOSITION

Technical Field

The present invention relates to a medical grade,
5 antimicrobial-containing adhesive particularly suited for use
in skin contact applications, such as with surgical drapes,
tapes and wound dressings. More particularly, the adhesive
compound is essentially a 100% solids or essentially
solventless hot melt adhesive incorporating an acrylic polymer
10 in conjunction with diiodomethyl-p-tolylsulfone as an
antimicrobial agent.

Background of the Invention

It is recognized that numerous pathogens are present on
human skin. Therefore, in a hospital environment, it is
15 generally desired that the growth of disease-producing
microorganisms be inhibited, and preferably that these
microorganisms be destroyed so as to control patient infection
and encourage wound healing. Under most circumstances, the
bacteria of normal skin cannot cause wound infections, but in
20 the presence of foreign materials or open wounds, the
pathogenic potential of these bacteria appears to be
considerably enhanced. Furthermore, the likelihood of
bacterial contamination is at a peak immediately preceding,
during, and following surgical procedures. Accordingly, to
25 prevent contamination, it is imperative that the skin be
effectively disinfected before a surgical incision is made and
during the entire surgical procedure.

In response to such concerns, many topical antimicrobial agents have been developed. These agents typically are in the form of preoperative skin preps, surgical scrub tissues, washes, wound cleaners, lotions and ointments. A recognized limitation to such topical applications are a short effective delivery time. Microorganisms that may have survived the initial application of such a topical antimicrobial agent can act as a seed, causing the pathogen population in some instances to regenerate or rise to their initial levels.

Thus, continuous application of an antimicrobial agent to the site is recognized as a means of inhibiting this increase in population.

It has been recognized that a continuous or longer lasting antimicrobial effect may be achieved by incorporating the antimicrobial agent into an adhesive layer or into a surgical incise drape material itself.

Berglund et al. (U.S. Patent No. 4,310,509) disclose that it is known to incorporate biologically active agents into adhesive layers on a substrate to provide continuous application of such agent to the body. Disclosed examples of known adhesives containing antimicrobial agents include U.S. Patent No. 2,137,169, wherein phenol, thymol, methanol, etc. are added to a starch adhesive; U.S. Patent No. 3,249,109 where benzocaine was added to a tacky gelatin; U.S. Patent No. 3,632,740 where a corticosteroid is added to an adhesive; U.S. Patent No. 3,734,097 where a microencapsulated anti-neoplastic

agent is added to an adhesive; U.S. Patent No. 4,073,291 where Tretinooin is added to an adhesive; U.S. Patent No. 3,769,071 where 5-fluorouracil is incorporated into an adhesive; and U.S. Patent No. 3,896,789 where retinoic acid is incorporated 5 into a pressure-sensitive adhesive tape. Berglund et al. further teach that the prior art attempts to include an antimicrobial agent in an adhesive did not include the use of a broad spectrum antimicrobial because such adhesives had been frustrated by uncontrollable release of the agent with 10 accompanying skin irritation in some patients, along with failure to obtain sufficient antimicrobial activity.

Berglund et al. disclose a pressure sensitive adhesive composition which contains chlorhexidene, polyvinylpyrrolidone iodine or iodine which is applied onto a polymer sheet 15 material, such as polyethylene or polyurethane, for use as a surgical drape. The disclosed drape is applied to the skin with the adhesive side contacting the skin so that the antimicrobial agent can be released from the adhesive to the wound area prior to and during incision. The process for 20 making the adhesive disclosed by Berglund et al. involves forming an emulsifiable concentrate or an organic solution concentrate of a broad spectrum antimicrobial agent and mixing it into an adhesive, such that the broad spectrum antimicrobial is homogeneously dispersed as a separate phase 25 throughout the adhesive medium. The homogenous dispersion is then spread or coated to a substantially uniform layer

followed by drying of the wet layer in order to remove the solvents.

Rosso et al. (U.S. Patent No. 4,323,557) disclose a drape incorporating a pressure sensitive adhesive utilizing n-vinylpyrrolidone residues in the polymer backbone. Iodine is complexed with these residues to provide an antimicrobial effect. Rosso et al. espouse the stability of the adhesive composition over the prior art compositions. By stable, Rosso et al. asserts that a composition coating of 11 grains per 24 sq. in. which is attached to a polyethylene sheet can be exposed to a temperature of 120°F and a relative humidity of 9% for two weeks or to a dose of 2.5 megarads of gamma irradiation without substantial alteration of the physical appearance or of the chemical activity as tested by the starch test and microbiological activity as tested by the zone inhibition assay. The disclosure of Rosso et al. is incorporated herein by reference.

The process for forming the adhesive composition disclosed by Rosso et al. involves forming a pressure-sensitive adhesive and mixing into it an antimicrobial treating solution comprising iodine, an iodide, and a solvent. The resulting composition preferably contains n-vinylpyrrolidone in the backbone of the pressure-sensitive adhesive which serves to complex the iodine. Rosso et al. disclose that the composition may be either attached directly onto a flexible backing substrate or formed onto a release

liner for later use. Once applied, the solvents are then evaporated by means known to the art, whereby an adhesive film is formed which is useable in or on tapes, drapes and other medical devices.

5 Mixon et al. (U.S. Patent No. 5,069,907) disclose a surgical drape having incorporated therein a broad spectrum antimicrobial agent. The drape comprises a synthetic polymeric film or fabric having incorporated therethrough an amount of antimicrobial agent. The drape may optionally have
10 an adhesive layer attached to one of its external surfaces, wherein the adhesive layer can have dispersed therethrough an antimicrobial agent. The preferred antimicrobial agent used is 5-chloro-2-(2,4-dichlorophenoxy)phenol. Suitable adhesives utilized include polyacrylate adhesives.

15 Mixon et al. disclose a large number of antimicrobial agents which were contemplated for use with the disclosed composition. These include metal salts, typical antibiotics, antibacterial agents such as chlorhexidine and its salts, quaternary ammonium compounds, iodophors such as povidone
20 iodine, acridine compounds, biguanidine compounds, and a preferred antimicrobial agent 5-chloro-2-(2,4-dichlorophenoxy)phenol. Mixon et al. further disclose that these same antimicrobial agents, which they propose to utilize within the polymer composition for their surgical drape, can
25 also be utilized in an adhesive composition. Mixon et al. further state that the antimicrobial agent can be directly

applied to the surgical drape in solution as an aqueous dispersion, as a hot melt, or by a transfer process using known techniques, such as knife, roller-coating, or curtain-coating methods. The transfer process is disclosed as 5 particularly preferred. In a transfer process, the adhesive emulsion, including water or a different solvent, optionally containing an antimicrobial agent, is spread onto a sheet of release paper and dried to remove the water or solvent. The surgical drape is then brought into contact with the adhesive 10 and calendered to insure that the adhesive adheres to the drape. The surgical drape will then generally include a release sheet covering the adhesive, and the release sheet on which the adhesive is deposited can be used for that purpose, or that release sheet can be removed and replaced with another 15 release sheet. In embodiments where the adhesive contains an antimicrobial agent, the mixture of adhesive and antimicrobial agent is dried after coating on the release sheet, and the antimicrobial agent remains dispersed in the adhesive.

Generally, presently known antimicrobial agents are 20 limited in their ability to withstand heat during processing. The lack of heat stability of n-vinyl pyrrolodione iodine has limited the ability for drapes having this antimicrobial agent from being ethylene oxide sterilized under heat stress. Further, many of the antimicrobial compounds cannot be 25 radiation sterilized. Thus, each prior art reference teaches that it is preferred to apply the antimicrobial adhesive in

conjunction with a solvent followed by subsequent evaporation of the solvent.

Accordingly, the need exists for an adhesive composition having an antimicrobial agent dispersed therethrough which is 5 heat stable and solventless. Such composition would eliminate the need for use of solvents with their potential environmental effects and would eliminate the need for removing such solvent from the adhesive after application to the drapes. Further, the heat stability would allow the 10 solventless adhesive to be applied in a hot melt process, while also allowing for ethylene oxide sterilization under heat stress or radiation sterilization.

Summary of the Invention

The present invention is an adhesive composition having 15 a broad spectrum antimicrobial agent dispersed therethrough. The adhesive composition is an essentially solventless composition (100% solids), which is heat stable so that it may be applied in a hot melt process, while also being capable of 20 ethylene oxide sterilization under heat stress without loss of effectiveness of the antimicrobial agent. Specifically, the adhesive is for skin-contact applications, for example, surgical drapes, tapes and wound dressings. The antimicrobial agent utilized is diiodomethyl-p-tolylsulfone with a preferred concentration of antimicrobial agents in the adhesive of about 25 0.1% to about 2% loading by weight.

The antimicrobial agent is homogeneously dispersed

through the adhesive layer. Active antimicrobial molecules continually disassociate from the surface or leach out of the adhesive matrix over time, delivering biocidal activity at a distance from the adhesive surface. Applicants have 5 conclusively demonstrated this property by zone of inhibition tests on a wide variety of infectious organisms. These tests conclusively showed that microbes were inhibited at a distance from the sample.

Adhesive compositions can incorporate acrylic or rubber 10 based polymers to form the hot melt adhesive. A preferred composition includes a mixture of two acrylic polymers, one of which is a low molecular weight solid acrylic polymer, the other a medium molecular weight solid acrylic polymer, which are both designed for hot melt pressure-sensitive adhesive 15 applications. A low molecular weight solid acrylic polymer is available from Schenectady International, Inc. as Product No. HRJ-4326, and a medium molecular weight solid acrylic polymer is also available from Schenectady International, Inc. under Product No. HRJ-10127. Tackifiers can also be added to the 20 adhesive composition as is well known in the art.

The present adhesive composition is a hot melt adhesive. By hot melt adhesive, it is meant that the adhesive is essentially solventless or 100% solids and is processed in liquid form at elevated temperatures in the range of about 25 275°F to 350°F. A preferred temperature range for compounding and coating the antimicrobial adhesive is 290°F to 320°F. The

antimicrobial containing adhesive is manufactured by heating
the adhesive composition to about 250°F, including both a low
molecular weight acrylic polymer and a medium molecular weight
acrylic polymer along with any tackifiers to be utilized. The
5 mixture is then heated to about 310°F to about 315°F and mixed
until uniform with subsequent cooling to 290°F to 295°F at
which point the diiodomethyl-p-tolylsulfone is added. The
composition is mixed until uniform with subsequent packaging
and cooling. The composition may then be hot melted and
10 applied as needed by the user.

In a preferred application, the antimicrobial adhesive
composition of the present invention is utilized to overly a
polymeric substrate to form a surgical drape. The polymeric
substrate is preferably a polyester or co-polyester sheet
15 material which may have incorporated therein or coated on the
side opposite the adhesive an antimicrobial agent.

These and various other advantages and features of
novelty which characterize the present invention are pointed
out with particularity in the claims annexed hereto and
20 forming a part hereof. However, for a better understanding of
the invention, its advantages, and the objects obtained by its
use, reference should be made to the accompanying descriptive
matter in which there are illustrated and described preferred
embodiments of the present invention.

25 Detailed Description of the Preferred Embodiments

As required, detailed embodiments of the present

invention are disclosed herein. However, it is to be understood that the disclosed embodiments are merely exemplary of the present invention which may be embodied in various systems. Therefore, specific details disclosed herein are not 5 to be interpreted as limiting, but rather as a basis for the claims and as a representative basis for teaching one skilled in the art to variously practice the present invention.

The present invention is an adhesive compound which incorporates an adhesive component together with a broad 10 spectrum antimicrobial agent dispersed therethrough. The antimicrobial agent is homogeneously dispersed throughout the adhesive layer. Active antimicrobial molecules of the present composition disassociate from the surface or leach out of the adhesive matrix over time, delivering biocidal activity at a 15 distance from the adhesive surface.¹² Applicants have conclusively demonstrated by zone of inhibition tests on a wide variety of infectious organisms the efficacy of the present composition. These tests showed that microbes were inhibited and killed at a distance from the sample as detailed 20 in the attached experimental examples.

The adhesive of the present invention is specifically suited for use in skin contact applications during and after medical procedures, for example, as an adhesive in surgical drapes, wound dressings and tapes. The adhesive composition 25 is a hot melt adhesive. By hot melt adhesive, it is meant that the adhesive is essentially solventless or 100% solids

and flowable at elevated temperatures for application to a substrate material, such as a surgical drape. The preferred adhesive composition incorporates acrylic polymers and added tackifiers to form a pressure-sensitive adhesive which is
5 particularly suited for use in surgical procedures.

A preferred combination of acrylic polymers to form the adhesive composition includes the combination of a low molecular weight solid acrylic polymer and a medium molecular weight solid acrylic polymer in a ratio of about 1 to 4,
10 respectively, to optimize the adhesion of the adhesive to skin, cohesion and resistance to cold flow. A low molecular acrylic polymer is a polymer having a molecular weight ranging from about 90,000 to about 120,000, while a medium molecular weight acrylic polymer has a molecular weight ranging from
15 about 140,000 to about 160,000. Suitable low molecular weight solid acrylic polymers and medium molecular weight solid acrylic polymers are available from Schenectady International, Inc. under Product Nos. HRJ-4326 and HRJ-10127, respectively.

The adhesive component of the composition can also
20 include tackifiers as are well known in the art. Tackifiers contemplated include SYLVATEC, ZONAREZ and FORAL which are available from Arizona Chemical and Hercules, Inc.

As previously stated, the adhesive compound is a hot melt adhesive. A preferred composition has a feasible temperature
25 range for working with the hot melt adhesive in the range of about 275°F to 350°F. The preferred temperature range for

compounding and coating with the adhesive is 290°F to 320°F.

Applicants have found that the addition of a heat stable antimicrobial agent to the above adhesive composition results in an effective antimicrobial adhesive composition. In 5 particular, Applicants have found that the addition of diiodomethyl-p-tolylsulfone to the above adhesive composition results in an effective antimicrobial adhesive which retains desirable properties during use and application at 275°F to about 350°F. A preferred loading of antimicrobial agent to 10 the adhesive is in the range of about 0.1% to about 2% by weight. A preferred loading is about 0.2% by weight to about 0.6% by weight of diiodomethyl-p-tolylsulfone to adhesive. The resulting heat stable antimicrobial containing adhesive is 15 100% solids and eliminates the need for use of a solvent and the requisite evaporation of such solvent. The hot melt adhesive can also be ethylene oxide sterilized under heat stress or radiation sterilized without loss of effectiveness of the antimicrobial.

A preferred source of diiodomethyl-p-tolylsulfone is 20 AMICAL 48, available from Angus Chemical Company.

The antimicrobial containing adhesive composition of the present invention is manufactured by mixing thoroughly at elevated temperature the acrylic polymers and tackifiers. A temperature of about 250°F to about 260°F has been found to be 25 adequate. Once mixed, the polymers and tackifiers are heated to 310°F to 350°F with continued mixing until uniform,

followed by cooling to 290°F to 295°F. The diiodomethyl-p-tolylsulfone is then added to the polymer and mixed until uniform. The resultant composition is packaged and cooled for subsequent hot melt applications.

5 As detailed below, the antimicrobial adhesive of the present invention was shown to be effective against a wide variety of microorganisms. The antimicrobial activity was determined by using a series of zone of inhibition tests, as are well known in the art. The effective release of
10 antimicrobial from the adhesive is estimated from the measurement of a zone of inhibition (an area of inoculated plate where organisms do not grow) surrounding the sample.

The adhesive utilized for the tests included 2% diiodomethyl-p-tolylsulfone homogeneously dispersed as
15 detailed above in an adhesive composition. The adhesive composition included 17% low molecular weight acrylic polymer (HRJ-4326 from Schenectady International, Inc.) and 67% medium molecular weight polymer (HRJ-10127 from Schenectady International, Inc.) along with 16% FLORAL 105 synthetic resin
20 from Hercules, Inc. as a tackifier. The adhesive composition was prepared as detailed above. The adhesive composition was then melted and applied to a substrate layer in a thin coating (approximately 0.05mm in thickness). The substrate was a co-polyester surgical drape material available from DuPont under
25 the tradename HYTREL. The coated substrate was cut to 6.0mm disks for use in testing.

Adhesive coated disks were then exposed to microorganisms using the following procedure:

1. A microbial suspension containing $\approx 1.0 \times 10^8$ organisms per ml in TSB was compared to the turbidity of a 0.5 MacFarland Standard.

2. A sterile swab was dipped into the culture suspension. The swab was rotated several times, pressing firmly on the inside wall of the tube above the fluid level. This removed excess inoculum from the swab.

10 3. The surface of a Mueller Hinton agar plate was inoculated by streaking the swab over the entire sterile agar surface. This streaking procedure was repeated two more times, rotating the plate approximately 60° each time to ensure an even distribution of inoculum.

15 4. The paper liner was removed from each 6mm adhesive coated disc and the film was aseptically placed adhesive side down on the surface of the inoculated agar plate. Control samples were handled identically.

20 5. Immediately following the addition of the discs, the Mueller Hinton agar plates were placed in ambient air at 35-37° for 18-24 hours. Following incubation, the zones of inhibition surrounding the discs were measured. When no zone was observed, the disc was aseptically removed and the area beneath the disc was evaluated for growth of the test 25 organism. The tests were repeated two or three times, using relevant microorganisms. Experimental results are presented

in the table below, reported as the average diameter zone of inhibition surrounding/under 6.0mm samples. A 6.0 mm zone of inhibition indicates no growth of the test organism beneath the 6.0 mm test discs, while larger zones indicate effective antimicrobial activity at a distance from the disc.

Table 1

	Test Organism	Zone of Inhibition
10	Staphylococcus aureus (ATCC 6538)	12.0mm
	Escherichia coli (ATCC 11229)	6.0mm
15	Pseudomonas aeruginosa (ATCC 15442)	6.0mm
	Klebsiella pneumoniae (ATCC 4352)	7.0mm
20	Pseudomonas cepacia (ATCC 25416)	6.0mm
	Enterobacter cloacae (ATCC 13047)	6.0mm
25	Serratia marcescens (ATCC 14746)	6.5mm
	Streptococcus pyogenes (ATCC 19615)	10.5mm
30	Enterococcus faecalis- Vancomycin Resistant (ATCC 51299)	9.5mm
	Candida albicans (ATCC 10231)	33.5mm
	Bacillus subtilis (ATCC 19659)	9.2mm

These results indicate that the present adhesive is effective in inhibiting these eleven relevant organisms, even

after hot melt processing and ethylene oxide sterilization.

New characteristics and advantages of the invention covered by this document have been set forth in the foregoing description. It will be understood, however, that this disclosure is, in many respects, only illustrative. Changes may be made in details, particularly in matters of shape, size, and arrangement of parts, without exceeding the scope of the invention. The scope of the invention is, of course, defined in the language in which the appended claims are expressed.

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